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NEW FUNCTIONAL ALLYLIC LITHIUM REAGENTS: GEM-DIALKOXYALLYLLITHIUM-ETC(U)  
FEB 80 D SEYFERTH, R E MAMMARELLA, H A KLEIN N00014-76-C-0837  
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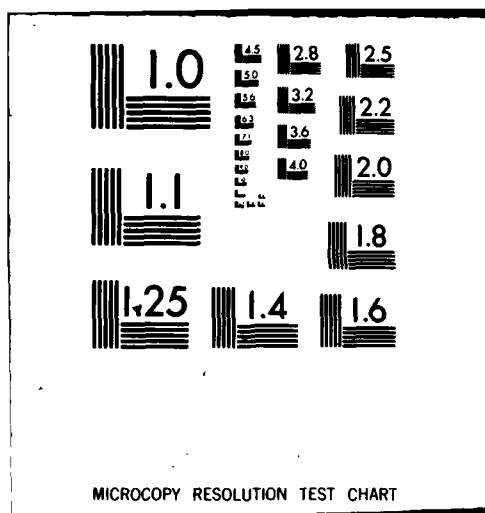
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20. with allyl bromide gave esters of 5-hexenoic acid,  
 $\text{CH}_2=\text{CH}(\text{CH}_2)_3\text{CO}_2\text{R}$  (R = Me, Et). ↗

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(6) NEW FUNCTIONAL ALLYLIC LITHIUM REAGENTS: GEM-DIALKOXYALLYLLITHIUM  
REAGENTS: A USEFUL ROUTE TO  $\alpha$ -SILYL AND  $\alpha$ -STANNYLPROPIONATE ESTERS.

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NEW FUNCTIONAL ALLYLIC LITHIUM REAGENTS: GEM-DIALKOXYALLYLLITHIUM  
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Dietmar Seydel\*, Robert E. Mammarella and Helmut A. Klein

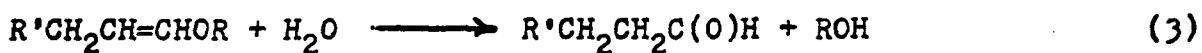
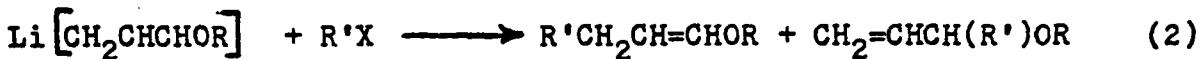
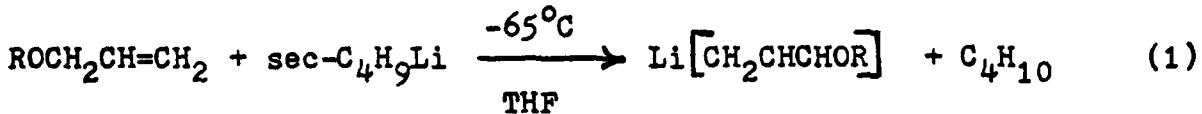
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ABSTRACT

The reaction of sec-butyllithium with acrolein dialkyl acetals in THF or in THF/Et<sub>2</sub>O/pentane at -95°C results in formation of gem-dialkoxyallyllithium reagents, Li[CH<sub>2</sub>CHC(OR)<sub>2</sub>]. These react with organosilicon and organotin chlorides to give ketene acetals, R<sub>3</sub>SiCH<sub>2</sub>CH=C(OR)<sub>2</sub> and R<sub>3</sub>SnCH<sub>2</sub>CH=C(OR)<sub>2</sub>. The acid hydrolysis of these products produces  $\beta$ -substituted propionic acid esters, R<sub>3</sub>SiCH<sub>2</sub>-CH<sub>2</sub>CO<sub>2</sub>R and R<sub>3</sub>SnCH<sub>2</sub>CH<sub>2</sub>CO<sub>2</sub>R. Reactions of these lithium reagents with allyl bromide gave esters of 5-hexenoic acid, CH<sub>2</sub>=CH(CH<sub>2</sub>)<sub>3</sub>-CO<sub>2</sub>R (R = Me, Et).

### INTRODUCTION

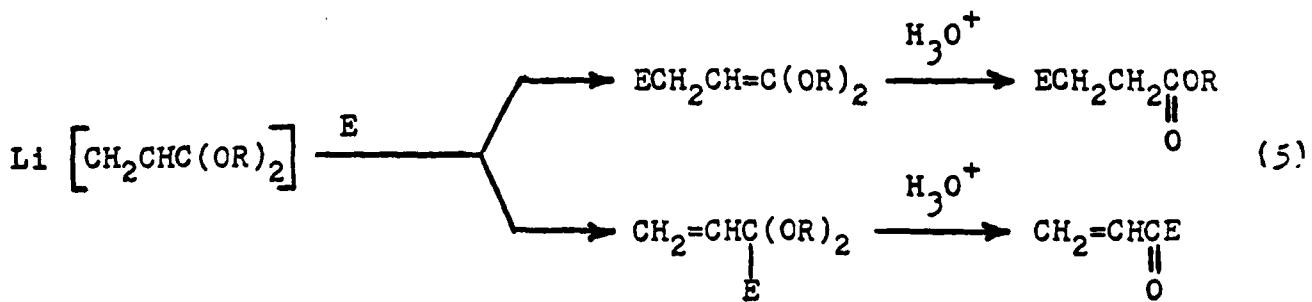
The lithiation of allylic ethers provides useful reagents which are operationally equivalent to a  $\beta$ -acyl carbanion (eq. 1-3)<sup>1</sup>.



In this context, the possible lithiation of an allylic acetal,  $\text{CH}_2=\text{CHCH}(\text{OR})_2$ , was of some interest. The expected product is a gem-dialkoxyallyllithium reagent (eq. 4) which potentially could react



with an electrophile, E, at either the unsubstituted or the substituted terminus of the allyl group (eq. 5). If the new bond is formed



at the substituted end, an allylic acetal would be formed. Hydrolysis of the latter would give a vinyl ketone. In this case the reagent would function as a carbonyl anion equivalent,  $\text{CH}_2=\text{CHC(O)}^-$ . If, on the other hand, the new bond is formed at the unsubstituted terminus of the allyl group, a  $\gamma$ -substituted ketene acetal would result. Hydrolysis of such a product would give an ester, as shown in eq. 5. In

in this case the reagent would be operationally equivalent to a  $\beta$ -alkoxycarbonyl carbanion,  $\text{ROC(O)CH}_2\text{CH}_2^-$ .

Of special interest to us were the reactions of such  $\text{Li}[\text{CH}_2\text{CHC(OR)}_2]$  reagents with organometallic halides, in particular trimethylchlorosilane and trimethyltin chloride. Alkyl- and aryl-substituted allylic lithium reagents of type  $\text{Li}[\text{CH}_2\text{CHCHR}]$  and  $\text{Li}[\text{CH}_2\text{CHCR}_2]$  had been found to react with both of these halides to give exclusively products of structure  $\text{Me}_3\text{MCH}_2\text{CH=CHR}$  and  $\text{Me}_3\text{MCH}_2\text{CH=CR}_2$  ( $\text{M} = \text{Si}$  and  $\text{Sn}$ ), respectively.<sup>2,3</sup> On the other hand, trimethylchlorosilane reacted with gem-dichloro-<sup>4</sup> and gem-difluoroallyllithium<sup>5</sup> to give  $\text{Me}_3\text{SiCX}_2\text{CH=CH}_2$  ( $\text{X} = \text{Cl}$  and  $\text{F}$ ). Trimethyltin chloride formed  $\text{Me}_3\text{SnCH}_2\text{CH=CCl}_2$  on reaction with  $\text{Li}[\text{CH}_2\text{CHCCl}_2]$ . In the case of the  $\text{Me}_3\text{SiCl}$  reactions it was quite likely that products of kinetic control were obtained, while those derived from  $\text{Me}_3\text{SnCl}$  were products of thermodynamic control.<sup>4</sup>

In view of the potential synthetic utility of reagents of  $\text{Li}[\text{CH}_2\text{CHC(OR)}_2]$  and in view of the uncertainty of the regioselectivity in their reactions with various electrophiles, we have investigated their synthesis, stability and, to a limited extent, their reactions.

#### RESULTS AND DISCUSSION

We have found that acrolein dimethyl and diethyl acetals may be lithiated by sec-butyllithium in a THF/diethyl ether/pentane solvent mixture at  $-90^\circ \pm 5^\circ\text{C}$  to give a yellow solution containing the respective gem-dialkoxyallyllithium reagents (eq. 4,  $\text{R} = \text{Me}$  and  $\text{Et}$ ;  $\text{R}' = \text{sec-C}_4\text{H}_9$ ). The temperature range for successful preparation of these reagents is quite narrow: below  $-100^\circ\text{C}$  the metalation reaction appears to be too slow; above  $-85^\circ\text{C}$  the reagents begin to decompose slowly. Their decomposition is rapid at  $-65^\circ\text{C}$  and above. Nonetheless-

TABLE 1. Reactions of gem-Dialkoxyallyllithium Reagents<sup>a</sup> with Chlorosilanes, Tin Chlorides and Allyl Bromide.

Acrolein Dialkyl Acetal (mmol)	Substrate (mmol)	Work-up	Product (% Yield) <sup>d</sup>
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 21 )	$\text{Me}_3\text{SiCl}$ ( 27.6 )	5% HCl	$\text{Me}_3\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Me}$ ( 75 )
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 42 )	$\text{Me}_3\text{SiCl}$ ( 45 )	anhydrous	$\text{Me}_3\text{SiCH}_2\text{CH}=\text{C}(\text{OMe})_2$ ( 70 ) <sup>b</sup>
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 43 )	$\text{Me}_2\text{SiCl}_2$ ( 21 )	5% HCl	$\text{Me}_2\text{Si}(\text{CH}_2\text{CH}_2\text{CO}_2\text{Me})_2$ ( 61 )
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 21 )	$\text{Et}_3\text{SiCl}$ ( 21 )	5% HCl	$\text{Et}_3\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Me}$ ( 70 ) <sup>c</sup>
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 42 )	$\text{PhMe}_2\text{SiCl}$ ( 49 )	5% HCl	$\text{PhMe}_2\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Me}$ ( ?? )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 25 )	$\text{Me}_3\text{SiCl}$ ( 40 )	$\text{H}_2\text{O}$	$\text{Me}_3\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 43 )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 63 )	$\text{Et}_3\text{SiCl}$ ( 86 )	$\text{H}_2\text{O}$	$\text{Et}_3\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 86 ) ( 71% distilled )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 63 )	$\text{PhMe}_2\text{SiCl}$ ( 86 )	$\text{H}_2\text{O}$	$\text{PhMe}_2\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 53 )
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 42 )	$\text{Me}_3\text{SnCl}$ ( 60 )	10% HCl	$\text{Me}_3\text{Sn}(\text{CH}_2\text{CH}_2\text{CO}_2\text{Me})$ ( 72 )
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 42 )	$\text{Me}_3\text{SnCl}$ ( 60 )	anhydrous	$\text{Me}_3\text{SnCH}_2\text{CH}=\text{C}(\text{OMe})_2$ ( 69 )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 25 )	$\text{Me}_3\text{SnCl}$ ( 40 )	5% HCl	$\text{Me}_3\text{SnCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 27 )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 25 )	$\text{Me}_3\text{SnCl}$ ( 40 )	$\text{H}_2\text{O}$	$\text{Me}_3\text{SnCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 1 part ) + $\text{Me}_3\text{SnCH}_2\text{CH}=\text{C}(\text{OEt})_2$ ( 2 parts )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 25 )	$\text{Et}_3\text{SnCl}$ ( 32 )	5% HCl	$\text{Et}_3\text{SnCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 24 )
$\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ( 42 )	$\text{CH}_2=\text{CHCH}_2\text{Br}$ ( 60 )	5% HCl	$\text{CH}_2=\text{CHCH}_2\text{CH}_2\text{CO}_2\text{Me}$ ( 47 )
$\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ( 25 )	$\text{CH}_2=\text{CHCH}_2\text{Br}$ ( 40 )	5% HCl	$\text{CH}_2=\text{CHCH}_2\text{CH}_2\text{CO}_2\text{Et}$ ( 22 )

TABLE 1: Footnotes

<sup>a</sup> Prepared at -90° 5°C by reaction with sec-butyllithium.

<sup>b</sup> A 12:1 mixture of  $\text{Me}_3\text{SiCH}_2\text{CH}=\text{C}(\text{OMe})_2$  and  $\text{Me}_3\text{SiCH}_2\text{CH}_2\text{CO}_2\text{Me}$  was produced.

<sup>c</sup> The  $\text{Et}_3\text{SiOSiEt}_3$  formed on hydrolysis of unconverted  $\text{Et}_3\text{SiCl}$  cannot be separated from the product and so it is better to use an excess of the lithium reagent.

<sup>d</sup> By-products included the respective disiloxane in those reactions in which an excess of the chlorosilane was used.

acetal. Organotin chlorides, on the other hand, do not hydrolyze readily. Although their rate of hydrolysis is rapid, the hydrolysis equilibrium is not favorable.<sup>6</sup> Hence, insufficient acid is generated and external acid must be added to effect hydrolysis of the acetal.

The formation of products of type  $R'^3MCH_2CH=C(OR)_2$  rather than  $R'^3MC(OR)_2CH=CH_2$  in these reactions of  $Li[CH_2CHC(OR)_2]$  with  $R'^3MCl$  may be understood in terms of the operation of steric effects. It is unlikely that allylic organosilicon compounds of either type ( $M = Si$  in the formulas above) would take part in transmetalation equilibria, so we assume that we are dealing with products of kinetic control. Products of type  $R'^3SiC(OR)_2CH=CH_2$  would have been of some interest: their hydrolysis should have given  $\alpha$ -silyl ketones,  $R'^3SiC(O)CH=CH_2$ .

Other methods for the preparation of  $\beta$ -silyl- and  $\beta$ -stannyl-propionic acid esters are available. Such silyl-substituted esters can be prepared by the acetoacetic ester and malonic ester syntheses when the appropriate halomethylsilanes,  $R_3SiCH_2X$ , are available.<sup>9</sup> Both silyl and stannyl-substituted esters have been prepared by silicon or tin hydride addition to acrylate esters,<sup>10-13</sup> while the reaction of  $\beta$ -halopropionic acid esters with tin foil provided another route to  $\beta$ -stannyl-substituted esters.<sup>14</sup> Thus the present procedure, which is based on the availability of the silicon or tin halide, complements these procedures.

EXPERIMENTAL

General Comments.

All reactions were carried out in flame-dried, nitrogen-flushed glassware under an atmosphere of prepurified nitrogen or argon. Solvents were rigorously dried prior to use. The reaction temperatures which are reported are uncorrected. They were obtained using a pentane total immersion thermometer which was immersed to a depth of about 3 cm into the stirred reaction mixture. Experiments showed that a stem correction of  $-8^{\circ}$  to  $-10^{\circ}\text{C}$  is appropriate.

sec-Butyllithium was purchased from Alfa Products, Thiokol/Ventron Corp., chlorosilanes from Petrarch Systems, Inc., the acrolein dialkyl acetals from Aldrich Chemical Co. Methyltin starting materials were kindly donated by Cincinnati Milacron Chemicals, Inc.

Infrared spectra were recorded using a Perkin Elmer Model 457A grating infrared spectrophotometer, NMR spectra with a Varian Associates T60 spectrometer. Gas-liquid chromatography (GLC) was employed for analysis of reaction products, isolation of product samples and for yield determination by the internal standard method.

Preparation of gem-Dialkoxallyllithium Reagents: General Procedure.

A Morton (creased) flask of appropriate size, equipped with a paddle-type stirrer, a no-air stopper and a Claisen adapter which was fitted with a low temperature thermometer and a gas inlet tube, was flame-dried and then allowed to cool under a stream of nitrogen or argon. The flask then was charged with the solvent mixture (75 ml of THF, 15 ml of diethyl ether and 15 ml of pentane), 21 mmol of the acrolein dialkyl acetal (2.5 ml in the case of  $\text{CH}_2=\text{CHCH}(\text{OMe})_2$ ) was added, and the solution was cooled to  $-90^\circ \pm 5^\circ\text{C}$  by partial immersion in a liquid nitrogen Dewar flask. To this solution was added with stirring and under nitrogen (or argon) 20 ml of 1.1M sec-butyl-lithium in cyclohexane (22 mmol). The temperature was maintained at  $-90^\circ \pm 5^\circ\text{C}$  during the course of this very slow, dropwise addition. The resulting yellow solution was stirred 2-3 hr. at  $-90^\circ\text{C}$ .

To this reagent solution then was added the chlorosilane or the tin chloride (slight excess, ca. 25-30 mmol) by syringe over a 10 min. period. The resulting mixture, now milky white, was stirred at  $-90^\circ\text{C}$  for 30 min. and then was allowed to warm to room temperature.

In the non-hydrolytic work-up, the reaction mixture was transferred by cannula to a dry distillation flask. Solvents were removed at reduced pressure and the residue was trap-to-trap distilled at ca. 0.1 mm Hg into a flask cooled with liquid nitrogen. The distillate then was examined by GLC.

In the hydrolytic work-up, the cannulated solution was poured into a separatory funnel and treated successively with distilled water, two portions of 5 or 10% hydrochloric acid and, again, water. The aqueous phases were back-extracted with pentane and the combined organic phases were dried ( $\text{MgSO}_4$ ) and the solvents removed at

reduced pressure. The residue was trap-to-trap distilled, as above.

Some reactions were carried out on a larger scale (42 mmol of  $\text{CH}_2=\text{CHCH}(\text{OMe})_2$ , 63 mmol of  $\text{CH}_2=\text{CHCH}(\text{OEt})_2$ ). In some cases THF alone, rather than the mixed solvent system, was used, but the mixed solvent is preferred.

Organosilicon and Organotin Products: Characterization.

Table 2 lists the organosilicon and organotin esters and acetals prepared, their refractive indices, analyses and spectro-

scopic properties. The infrared spectra of the esters (liquid film) showed strong ester carbonyl absorption at  $1735-1740 \text{ cm}^{-1}$ . The ketene acetals showed strong bands in their IR spectra at 1675 ( $\text{Me}_3\text{SiCH}_2\text{CH}=\text{C}(\text{OMe})_2$ ) and  $1670 \text{ cm}^{-1}$  ( $\text{Me}_3\text{SnCH}_2\text{CH}=\text{C}(\text{OMe})_2$ ).

The methyl and ethyl esters of 5-hexenoic acid are known compounds.

$\text{CH}_2=\text{CHCH}_2\text{CH}_2\text{CH}_2\text{CO}_2\text{Me}$ ,  $n^{20}\text{D}$  1.4205, a known compound.<sup>7</sup>

IR (film):  $\nu(\text{C}=\text{O})$  1755,  $\nu(\text{C}=\text{C})$   $1640 \text{ cm}^{-1}$ .

NMR ( $\text{CDCl}_3$ ):  $\delta$  1.6-2.5 (m, 6H,  $\text{CH}_2\text{CH}_2\text{CH}_2$ ), 3.6 (s, 3H,  $\text{CH}_3$ ) and 4.75-6.3 (m, 3H,  $\text{CH}=\text{CH}_2$ ).

$\text{CH}_2=\text{CHCH}_2\text{CH}_2\text{CH}_2\text{CO}_2\text{Et}$ ,  $n^{20}\text{D}$  1.4225; lit.<sup>8</sup>  $n^{20}\text{D}$  1.4220

IR (film):  $\nu(\text{C}=\text{O})$  1735,  $\nu(\text{C}=\text{C})$   $1640 \text{ cm}^{-1}$ .

NMR ( $\text{CCl}_4$ ):  $\delta$  1.22 (t, 3H, J 7Hz,  $\text{CH}_3$ ), 1.48-2.03 (m, 2H,  $\text{CH}_2\text{CH}_2\text{CH}_2$  2.10 and 2.22 (2t, 4H, J 7Hz,  $\text{CH}_2\text{CH}_2\text{CH}_2$ ); 4.03 (q, 2H,  $\text{OCH}_2$ ), 4.73-6.17 (m, 3H,  $\text{CH}=\text{CH}_2$ ).

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TABLE 1. Reaction Products.

Compound	$n^t_D$ ( $^9C$ )	Analysis, %		$^1H$ NMR $\delta$ (ppm)
		Found (Calcd.)	Carbon Hydrogen	
$Me_3SiCH_2CH_2CO_2CH_3$	1.4190 (20) <sup>a</sup>	52.42 (52.45)	10.04 (10.06)	0.00 (s, 9H, $Me_3Si$ ), 0.63-1.00 (m, 2H, $SiCH_2$ ), 2.1-2.44 (m, 2H, $CH_2CO$ ), 3.6 (s, 3H, $OCH_3$ ) (in $CDCl_3$ )
$Me_2Si(CH_2CH_2CO_2Me)_2$	1.4396 (25)	51.25 (51.22)	8.60 (8.62)	0.00 (s, 6H, $Me_2Si$ ), 0.65-1.05 (m, 4H, $SiCH_2$ ), 2.1-2.5 (m, 4H, $CH_2CO$ ), 3.6 (s, 6H, $OCH_3$ ) (in $CDCl_3$ )
$Et_3SiCH_2CH_2CO_2CH_3$	1.4409 (25)	59.19 (59.40)	10.67 (10.89)	0.15-1.25 (m, 17H, $Et_3Si$ and $SiCH_2$ ), 1.95-2.4 (m, 2H, $CH_2CO$ ), 3.55 (s, 3H, $OCH_3$ ) (in $CDCl_3$ )
$PhMe_2SiCH_2CH_2CO_2CH_3$	1.4950 (25)	64.74 (64.86)	8.21 (8.11)	0.2 (s, 6H, $SiMe_2$ ), 0.8-1.2 (m, 2H, $SiCH_2$ ), 2.0-2.4 (m, 2H, $CH_2CO$ ), 3.45 (s, 3H, $OCH_3$ ), 7.2-7.5 (m, 5H, Ph) (in $CDCl_3$ )
$Me_3SiCH_2CH_2CO_2C_2H_5$	<sup>b</sup>	54.91 (55.12)	10.40 (10.41)	0.00 (s, 9H, $Me_3Si$ ), 0.62-1.02 (m, 2H, $SiCH_2$ ), 1.23 (t, $J$ 7Hz, 3H, $CH_3$ of Et), 2.02-2.37 (m, 2H, $CH_2CO$ ), 4.02 (q, $J$ 7Hz, $CH_2$ of Et) (in $CCl_4$ )
$Et_3SiCH_2CH_2CO_2C_2H_5$	1.4407 (25) <sup>c</sup>	61.04 (61.05)	11.12 (11.18)	0.27-1.18 (m, 17H, $Et_3Si$ and $SiCH_2$ ), 1.27 (t, $J$ 7Hz, 3H, $CH_3$ of OEt), 2.03-2.38 (m, 2H, $CH_2CO$ ), 4.05 (q, 1

$\delta$  7Hz, 2H, OCH<sub>2</sub>)

PhMe <sub>2</sub> SiCH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> C <sub>2</sub> H <sub>5</sub>	1.4917 (25) <sup>d</sup>	66.09 (66.05)	8.52 (8.53)	0.32 (s, 6H, Me <sub>2</sub> Si), 1.05 (t, J 7Hz, 2H, SiCH <sub>2</sub> ), 1.33 (t, J 7Hz, 3H, CH <sub>3</sub> of OEt), 2.08-2.42 (m, 2H, CH <sub>2</sub> CO), 4.03 (q, 2H, CH <sub>2</sub> of OEt), 6.88-7.57 (m, 5H, Ph) (in CCl <sub>4</sub> )
Me <sub>3</sub> SnCH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> CH <sub>3</sub>	1.4695 (20) <sup>e</sup>	33.44 (33.50)	6.49 (6.38)	0.07 (s, 9H, Me <sub>3</sub> Sn, J(119Sn- <sup>1</sup> H) 52 Hz, J(117 <sub>H</sub> - <sup>1</sup> H) 50 Hz), 1.00 (t, J 8Hz, 2H, SnCH <sub>2</sub> ), 2.50 (t, J 8Hz, 2H, CH <sub>2</sub> CO), 3.63 (s, 3H, OCH <sub>3</sub> ) (in CDCl <sub>3</sub> )
Me <sub>3</sub> SnCH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> C <sub>2</sub> H <sub>5</sub>	1.4638 (25) <sup>f</sup>	36.64 (36.27)	6.92 (6.85)	0.08 (s, 9H, Me <sub>3</sub> Sn, J(119Sn- <sup>1</sup> H) 52 Hz, J(117 <sub>H</sub> - <sup>1</sup> H) 50 Hz), 0.95 (t, 7Hz, 2H, SnCH <sub>2</sub> ), 1.27 (t, J 7Hz, 3H, CH <sub>3</sub> of OEt), 2.50 (t, J 7Hz, 2H, CH <sub>2</sub> CO), 4.08 (q, 2H, OCH <sub>2</sub> ) (in CCl <sub>4</sub> ) , SnCH <sub>2</sub>
Et <sub>3</sub> SnCH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> C <sub>2</sub> H <sub>5</sub>	1.4760 (25) <sup>g</sup>	43.51 (43.04)	7.90 (7.88)	0.33-1.48 (m, 20H, Et <sub>3</sub> Sn and CH <sub>3</sub> of OEt), 2.50 (t, 7Hz, CH <sub>2</sub> CO), 4.08 (q, J 7Hz, 2H, OCH <sub>2</sub> ) (in CCl <sub>4</sub> )
Me <sub>3</sub> SiCH <sub>2</sub> CH=C(OMe) <sub>2</sub>	1.4354 (20)	55.15 (55.17)	10.70	0.00 (s, 9H, Me <sub>3</sub> Si), 1.28 (d, J 8Hz, 2H, SiCH <sub>2</sub> ), 3.13-3.81 (m, 1H, =CH), 3.50 and 3.56 (two s, each 3H, OCH <sub>3</sub> ) (in CDCl <sub>3</sub> )

$\text{Me}_3\text{SnCH}_2\text{CH}=\text{C}(\text{OMe})_2$  1.4845 (20) 36.15 (6.82) 0.05 (s, 9H,  $\text{Me}_3\text{Sn}$ ,  $J(119\text{Sn}-^1\text{H})$  52 Hz,  $J(117\text{Sn}-^1\text{H})$  50 Hz), 1.52 (d, J 8 Hz, 2H,  $\text{SnCH}_2$ ), 3.15-3.90 (m, 1H, =CH), 3.46 and 3.53 (two s, each 3H,  $\text{OCH}_3$ ) (in  $\text{CDCl}_3$ )

$\text{Me}_3\text{SnCH}_2\text{CH}=\text{C}(\text{OEt})_2$  1.4640 (25) 40.90 (41.00) 7.65 (7.57) 0.07 (s, 9H,  $\text{Me}_3\text{Sn}$ ,  $J(119\text{Sn}-^1\text{H})$  52 Hz,  $J(117\text{Sn}-^1\text{H})$  50 Hz), 1.25 and 1.28 (two t, J 7Hz, each 3H,  $\text{CH}_3$  of OEt), 1.52 (d, J 8Hz, 2H,  $\text{SnCH}_2$ ), 3.42 (m, 5H,  $\text{OCH}_2$  and =CH) (in  $\text{CCl}_4$ )

a Lit. 15  $n^{20}\text{D}$  1.4180; b Lit. 9  $n^{20}\text{D}$  1.4198; lit. 16  $n^{20}\text{D}$  1.4205; c Lit. 12  $n^{20}\text{D}$  1.4422; d lit. 9

$n^{20}\text{D}$  1.4972; e Lit. 14  $n^{20}\text{D}$  1.4690; f Lit. 14  $n^{20}\text{D}$  1.4660; g Lit. 12  $n^{20}\text{D}$  1.4762.

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